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Article

Biochars as Innovative Humidity Sensing Materials

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Abstract: In this work, biochar-based humidity sensors were prepared by drop-coating technique. Polyvinylpyrrolidone (PVP) was added as an organic binder to improve the adhesion of the sensing material onto ceramic substrates having platinum electrodes. Two biochars obtained from different precursors were used. The sensors were tested toward relative humidity (RH) at room temperature and showed a response starting around 5 RH%, varying the impedance of 2 orders of magnitude after exposure to almost 100% relative humidity. In both cases, biochar materials are behaving as p-type semiconductors under low amounts of humidity. On the contrary, for higher RH values, the impedance decreased due to water molecules adsorption. When PVP is added to SWP700 biochar, n-p heterojunctions are formed between the two semiconductors, leading to a higher sensitivity at low RH values for the sensors SWP700-10% PVP and SWP700-20% PVP with respect to pure SWP700 sensor. Finally, response and recovery times were both reasonably fast (in the order of 1 min).

Keywords: humidity sensor; biochar; polyvinylpyrrolidone; n-p heterojunction

1. Introduction

Recent advances in science and technology have increased the demand for highly accurate, durable, sustainable, reproducible, and cheap humidity sensors. These sensors find many industrial applications ranging from environmental and meteorological monitoring, soil water content determination in agriculture, air conditioning systems, food quality monitoring, medical equipment, and other fields [1,2]. Humidity sensors can be classified as capacitive, resistive, or thermal conductive ones [3]. Usually, humidity sensors are based on metal oxide semiconductors, conducting polymers, carbon nanotubes, and/or graphene oxides. At room temperature, the adsorption of water vapor/molecules changes the resistance or the capacitance of the sensing film [4–15].

Carbon-based humidity sensors have attracted great attention since a couple of decades because of their large sensing area and high chemical inertness [16–20]. Biomass is a qualified carbon source, it is available in high quality and huge amount, and it is considered as an environmental friendly renewable resource [21]. In recent years, biochar applications invested in many fields [22], though the main application of this material remains field amendment in agriculture [22,23]. In addition, different biochars (residues of biomass pyrolysis) are becoming available from pilot plants producing biogas and energy [24,25]. Besides, in the last few years, biochar has been extensively studied as a substitute for more expensive carbon materials like carbon nanotubes, graphene, and others.

In a previous work, pyrolyzed bamboo was proposed as a sensing material with promising results, showing an excellent response toward humidity starting from 10% relative humidity (RH) [26]. However, the adhesion onto alumina substrates was minimal, and these films could be easily damaged

when testing, preventing their use in commercial products. Thus, in this research, the room temperature humidity sensing properties of two commercial biochars derived from different precursors [24,25] were investigated. First, the biochars were drop coated onto commercial alumina substrates with platinum interdigitated electrodes. Then, polyvinylpyrrolidone (PVP), acting as an organic binder was added to improve the adhesion onto the ceramic substrates. This polymer is used in many industrial applications, such as adhesives to enhance strength and toughness, paper manufacture for increasing strength and as a coating resin, as well as in synthetic fibers to accentuate dye receptivity. PVP polymers are also widely employed in inks, imaging, lithography, detergents, and soaps, in textile, ceramic, electrical, and metallurgical industries, and as a polymerization additive. Only in few studies was PVP used as humidity sensor in composites [27,28]. The sensing materials and the films were studied by means of Field Emission-Scanning Electron Microscopy (FESEM), X-Ray diffraction (XRD), Raman spectroscopy, X-ray photoelectron spectroscopy (XPS), and laser-granulometry measurements. The sensors were tested toward relative humidity at room temperature.

2. Materials and Methods

2.1. Materials

Two types of certified biochar were purchased from the UK Biochar Research Centre (BRC) (Edinburgh, UK) and used as innovative humidity sensing materials. The two biochars were selected among many available from BRC, based on the highest surface area and on the highest electrical conductivity. SWP700 is made from pyrolyzed mixed softwood pellets, obtained at a maximum treatment temperature of 700 °C with a heating rate of 87 °C/min. Its biochar yield is of 17.34%. OSR700 is produced from oil seed rape, by a pyrolysis process at 700 °C and a heating rate of 103 °C/min. In this case, a biochar yield of 22.62% is achieved. Table 1 reports the specific surface area (SSA) and the conductivity of both materials, as reported on their datasheets [24,25].

Table 1. Physical properties of OSR700 and SWP 700 biochars.

Biochar	Specific Surface Area (m ² /g)	Conductivity (S/m)
OSR700	25.2	0.311
SWP700	162.3	0.016

Both biochars in the form of pellets were first mixed in a mechanical grinder for 10 min and later manually ground in order to attain a homogeneous powder.

Polyvinylpyrrolidone (PVP, average Mw~13,000,000 by LS, Sigma Aldrich, Milano, Italy) was used as a polymeric binder and mixed with biochars in ethanol (for residual analysis, Sigma Aldrich ≥ 99.8% v/v).

2.2. Characterization of Biochars

SWP700 and OSR700 biochar powders were characterized by laser granulometry, X-Ray Diffraction (XRD), Raman Spectroscopy, X-Ray Photoelectron Spectroscopy (XPS), and Field Emission Scanning Electron Microscopy (FESEM) observations.

Particle size distribution was evaluated by laser granulometry (Mastersizer 3000, Malvern, Worcestershire, UK). The measurements were performed after manual grinding of the powders by means of an agate mortar and an agate pestle and sieving with a sieve having apertures of 45 µm. Laser granulometry measurements were carried out after dispersion in ethanol (500 ppm in volume) and sonication for 5 min.

XRD analysis was carried out with the aim of evidencing any possible crystallization peak among OSR700 and SWP700 raw materials. Spectra were recorded on a Pan'Analytical X'Pert Pro instrument (Pan'Analytical, Almelo, The Netherlands) working with Cu Kα radiation (0.154056 nm) in the range 5–70° 2θ, with a step size of 0.05° in 2θ and an acquisition time per step of 5 s. Diffraction patterns

were indexed by means of the Powder Data File database (P.D.F. 2000, International Centre of Diffraction Data, Newtown Square, PA, USA).

A Renishaw micro Raman (Renishaw, New Mills, UK) equipped with a green laser (514 nm) and a 50× objective was used to study the disorder and graphitization grade of the biochars under test. Raman characterization were performed at room temperature. The samples, in powder form, were exposed, without any pretreatment, to the micro laser beam (2 µm in diameter) for 10 s.

X-ray photoelectron spectroscopy (XPS) investigations were carried out by means of a PHI 5000 Versaprobe (Physical Electronics, Chanhassen, MN, USA) scanning X-ray photoelectron spectrometer (monochromatic Al K-alpha X-ray source with 1486.6 eV energy, 15 kV voltage, and 1 mA anode current) to investigate surface chemical composition.

Finally, morphologies of both kinds of powders and sensors were examined by means of an FESEM (Zeiss Supra-40, Oberkochen, Germany) equipped with an Oxford Energy Dispersive X-ray detector after the Cr sputtering process.

2.3. Sensor Fabrication

Platinum (5545-LS, ESL, King of Prussia, PA, USA) interdigitated electrodes were screen-printed on commercial α -alumina substrates (Coors Tek, Golden, CO, USA, ADS-96, 96% alumina, 0.85 cm × 1.7 cm). These electrodes were then fired at 980 °C (2 °C/min heating and cooling ramps) for 18 min, obtaining a high-grade adhesion and optimizing their electrical conductivity, as per the ink manufacturer's recommendations. Electrodes present a thickness of 400 µm, and their spacing is equal to 450 µm. In addition, the edge of an electrode belonging to one comb and the vertical connection of the second comb are separated by 400 µm.

To obtain biochar, PVP and biochar-PVP sensors, a drop coating technique was adopted by mixing with a magnetic stirrer 0.1 g of material (SWP700, OSR700, PVP, or biochar-PVP, respectively, in four different preparations) with 10 mL of ethanol in a beaker. After 1 h of mixing, solvent was evaporated at 80 °C, and sensors were realized by drop coating technique with a volume of 40 µL of this suspension onto α -alumina substrates with Pt electrodes by means of a micropipette.

Sensors were finally dried overnight at 80 °C (3 mg of material deposited) (Figure 1).

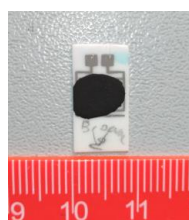


Figure 1. Image of a biochar sensor prepared by drop coating technique.

Subsequently, the sensors were tested toward relative humidity and cross-sensitivity tests were carried out toward ozone, ammonia, carbon dioxide, and methane at room temperature (SIAD, Bergamo, Italy; research quality gases).

2.4. Electrical Characterization

Humidity sensors were inserted into a laboratory apparatus made of a hermetic chamber in which relative humidity could be varied between 0 and 99% by steps, each one of 3 min. In fact, in this system, compressed air is separated into two flows: one is dehydrated over a chromatography alumina bed, while the second one is directed through two water bubblers, generating, respectively, a dry and a humid flow. Two precision microvalves recombine the two fluxes into one by means of a mixer and adjust the RH content [4]. All humidity measurements were carried out under a constant air flow of 1700 sccm (standard cubic centimeters). A commercial humidity and temperature probe was used as reference for temperature and RH values (Delta Ohm DO9406, Caselle di Selvazzano

(PD), Italy). All the measurements were done at room temperature (25 ± 0.5 °C). Relative humidity is at all temperatures and pressures defined as the ratio of the water vapor pressure (P_w) to the saturation water vapor pressure (over water, P_{ws}) at the gas temperature (T), according to Equation (1) and Ref. [29].

$$RH\% = \frac{P_w}{P_{ws}} \cdot 100 \quad (1)$$

The water vapor saturation pressure (in hPa) over water in the temperature range -20 °C– 50 °C is given by Equation (2).

$$P_{ws} = 6.116441 \times 10^{\left(\frac{7.591386 \cdot T}{T + 240.7263}\right)} \quad (2)$$

Thus, the water vapor pressure (in hPa) can be determined as follows (Equation (3)):

$$P_w = \frac{RH\% \cdot P_{ws}}{100} \quad (3)$$

During tests in a dynamic flow under variable amounts of relative humidity (RH), the sensors impedances were determined by means of a LCR meter (Hioki 3533-01, Nagano, Japan). The sensors were alimented by an AC voltage of 1 V at 1 kHz. The sensor response (SR), expressed in %, is defined as the relative variation of the starting impedance under dry air (Z_0), compared with the impedance measured under gas/humidity (Z_g) exposure (Equation (4)).

$$SR(\%) = 100 \times \frac{(Z_0 - Z_g)}{Z_0} \quad (4)$$

Moreover, cross-sensitivity measurements were carried out toward NH_3 (50 ppm in air), CO_2 (500 and 600 ppm in dry air and in air with 40% of relative humidity), CH_4 (100 ppm in air), and ozone (0.5 ppm in air) at room temperature under a constant flow of 1000 sccm, and sensors were put in another chamber of 0.25 L volume (in fact a three-neck glass flask). Target gas was diluted with air by means of flow meters (Teledyne Hastings Instruments HFM 300 controller and flow meters HFC 302, Teledyne Hastings, Hampton, VA, USA). Ozone was generated by a UV lamp (SOG-01, UVP-LLC Cambridge, UK) from a constant air flow of compressed air (1000 sccm). Ozone concentration was determined by the length of lamp exposed, according to calibration curves given by the lamp manufacturer. When performing measurements under both gas and humidity, a water bubbler was added to the flow of synthetic air used for the dilution of target gas.

The response times (the time taken by a sensor to achieve 90% of the total impedance change in the case of gas adsorption) together with the recovery times (the time necessary to reach 90% of the total impedance changes in the case of gas desorption) were also determined in the present work.

3. Results

3.1. Laser Granulometry

Laser granulometry evidenced that OSR700 biochar exhibits a bi-modal distribution, with maxima at 8 and 20 μm , and the mean diameter is equal to 10.5 μm , while SWP700 present a tri-modal distribution, with a mean diameter equal to 18.3 μm . These results are summarized in Figure 2 and Table 2.

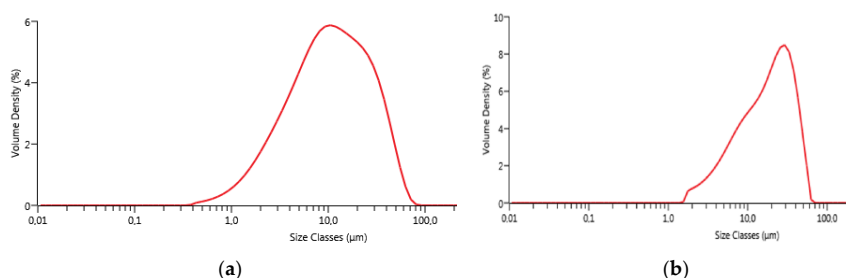


Figure 2. Laser granulometry measurement results: (a) OSR700; (b) SWP 700.

Table 2. Cumulative particle size distribution at 10% (D10), 50% (D50), and 90% (D90) for OSR700 and SWP 700 biochars.

Biochar	D10 (μm)	D50 (μm)	D90 (μm)
OSR700	2.61	10.5	33.4
SWP700	4.98	18.3	39.8

3.2. XRD Analysis

XRD patterns in Figure 3 reveal the amorphous nature of SWP700, while OSR700 presents some quartz content (peaks at 20.87° , 26.65° , 29.43° , 20.26° , 59.96° and 67.97° 2θ), JCPDS card No. 461045. The peak at 29.43° in 2θ was not indexed.

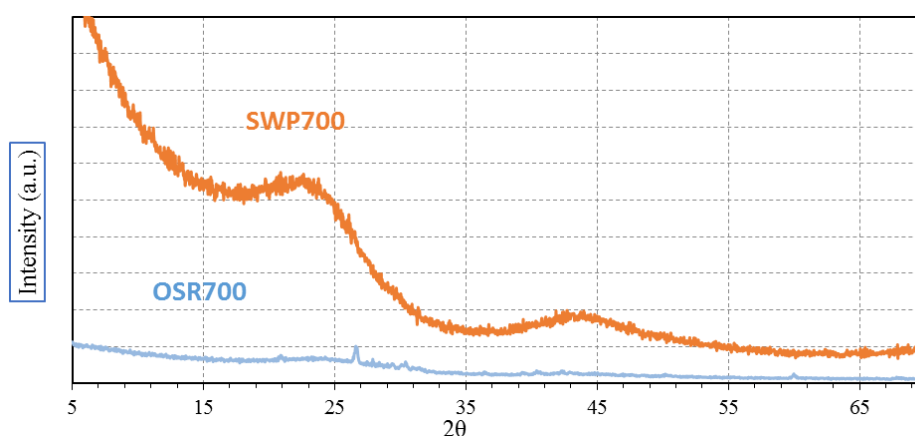


Figure 3. X-Ray diffraction (XRD), patterns of OSR700 and SWP700 biochars.

3.3. XPS Analysis

Surface chemistry of OSR700 and SWP700 were investigated using the XPS technique. The high-resolution spectra of C1s peaks are reported in Figure 4. XPS characterization shows the presence of the elements reported in Table 3. In particular, Si and Cl are present only in OSR700 while the carbon content is higher in SWP700. In both biochars, surface composition appears to be enriched in oxygen content compared to the information reported in the datasheets [24,25].

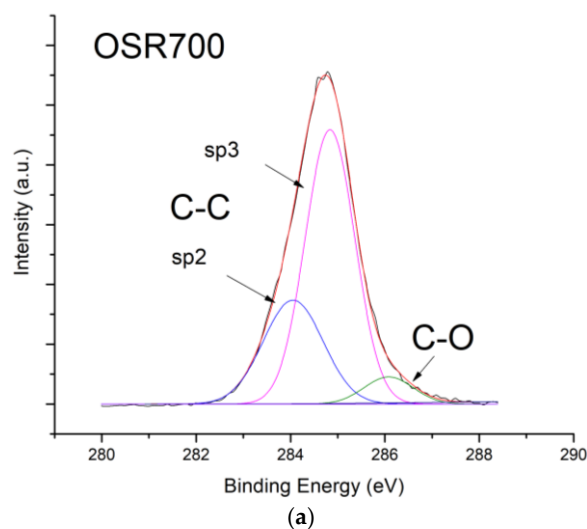


Figure 4. Cont.

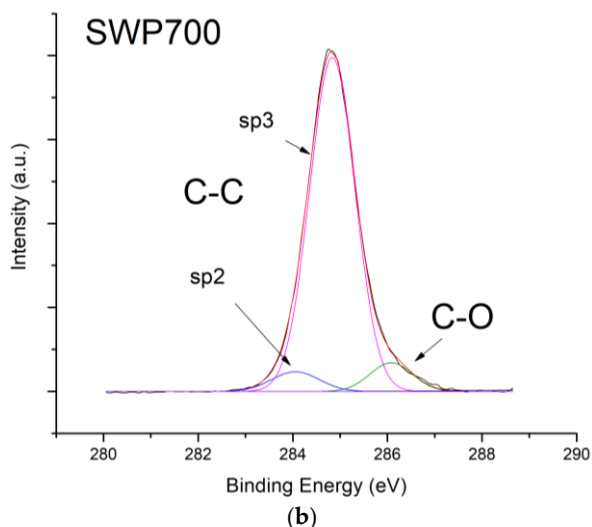


Figure 4. High resolution X-ray photoelectron spectroscopy (XPS) spectra of C1s peak for (a) OSR700; (b) SWP700.

Table 3. Weight% resulted from XPS characterization.

Weight%	OSR700	SWP700
C1s	55.8	72.1
C sp2	29.11	5.35
C sp3	63.68	87.13
C=O	6.21	6.96
O1s	23.3	16.1
K2s	15.2	10.3
Ca2p	3.2	1.2
Si2p	1.9	-
Cl2p	traces	-
N1s	traces	traces

(NB: all the elements were quantified from a survey, while C sp2, C sp3, and C=O content were determined from deconvolution of C 1s peaks in Figure 4).

OSR700 biochar presents a higher content of C in form of sp2 bonds compared to SWP700 (29.11% vs. 5.35% of total carbon). In addition, the concentration of other elements like K and Ca in OSR700 is higher, and Si is found only in the oil seed rape sample.

3.4. Raman Characterization

In the Raman investigation, different regions of the samples were analyzed to ensure the consistency of the results, that are depicted in Figure 5.

D ($\sim 1360\text{ cm}^{-1}$) and G ($\sim 1650\text{ cm}^{-1}$) peaks were focused. The signals were analyzed with inhouse developed and validated software that allows us (i) to get rid of the background contribution and (ii) to identify the contribution of the various components. The parameter focused in this work is the ratio I_D/I_G of the intensities of D and G peaks. 2D region, from 2300 to 3500 cm^{-1} , is also informative [30,31], but its investigation is beyond our interest in the present paper.

I_D/I_G ratio values are typical of carbonaceous material where the presence of defects is evident [30,31]. I_D/I_G ratios for the two biochars are comparable. Hence Raman analysis did not evidence remarkable dissimilarities between biochars, and it revealed a similar degree in structural defects.

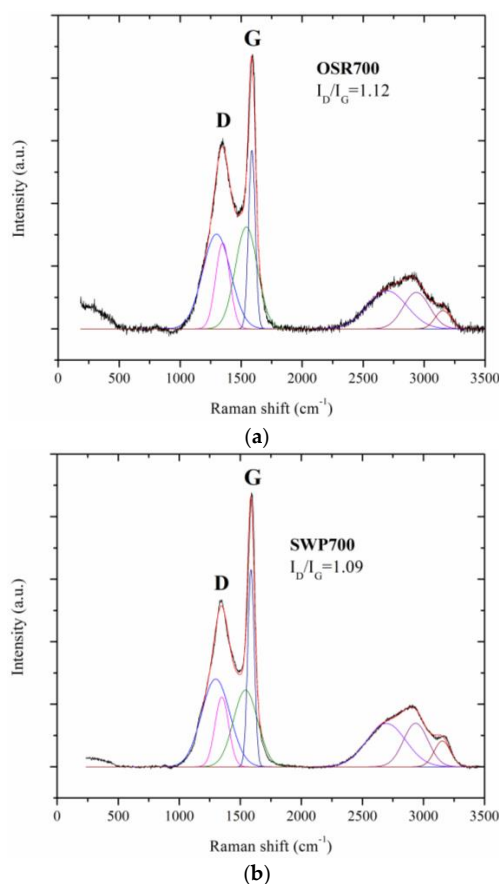


Figure 5. Raman spectra for (a) OSR700; (b) SWP700.

3.5. FESEM Observation

FESEM images in Figure 6 illustrate OSR700 biochar films at different magnification (2500 \times and 10,000 \times). A wide particle size distribution was observed, with some large, faceted particles surrounded by a finer fraction, in agreement with laser granulometry measurements. SWP700 biochar film observations are displayed in Figure 7, at 2500 \times and 10,000 \times magnification. The grinding process results in small fragments in the range between 1 and 50 μm , which is still in agreement with the results of particle size distribution measurements. The particles are faceted with sharp edges, as expected in a fragile powder exhibiting an irregular surface after grinding.

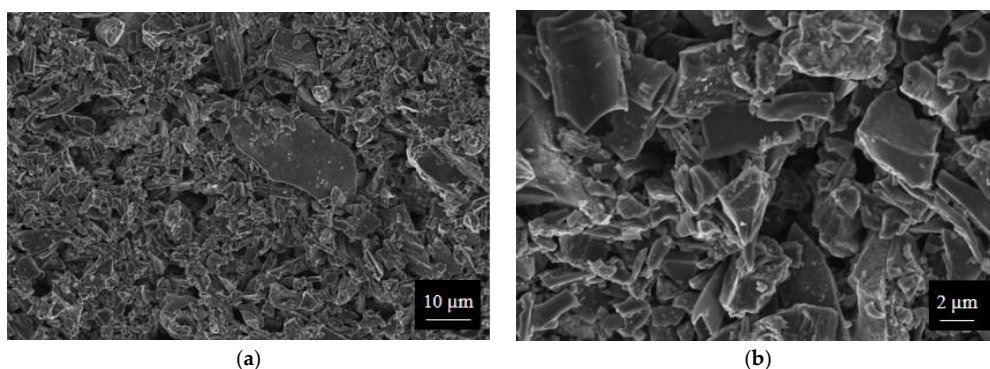


Figure 6. Field Emission-Scanning Electron Microscopy (FESEM) micrographs of OSR700 biochar sensor at different magnifications: (a) 2500 \times ; (b) 10,000 \times .

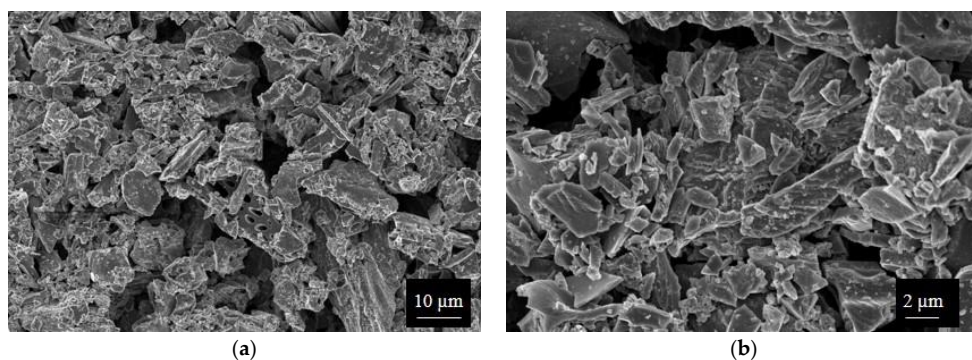


Figure 7. FESEM micrographs of SWP700 biochar sensor at different magnifications: (a) 2500 \times ; (b) 10,000 \times .

SWP700-PVP10% and SWP700-PVP20% film micrographs are illustrated in Figures 8 and 9. In these micrographs it is possible to observe an enhanced connection between biochar grains due to PVP coating.

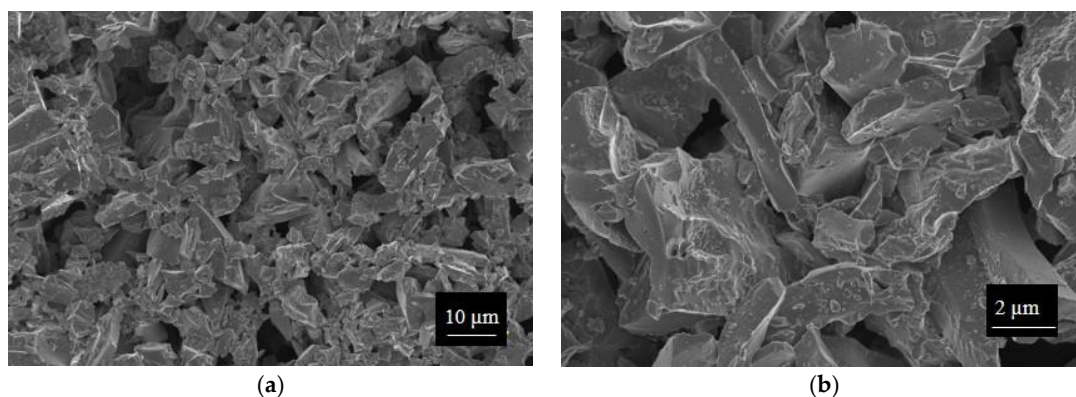


Figure 8. FESEM micrographs of SWP700-PVP10% sensor at different magnifications: (a) 2500 \times ; (b) 10,000 \times .

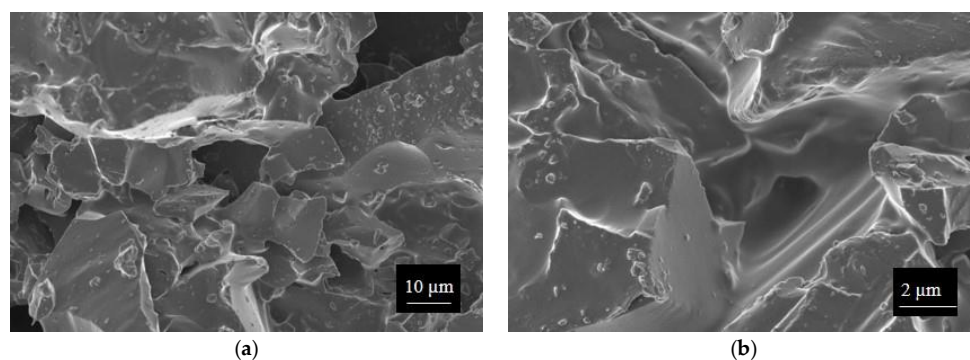


Figure 9. FESEM micrographs of SWP700-PVP20% sensor at different magnifications: (a) 2500 \times ; (b) 10,000 \times .

3.6. Electrical Characterization

Results of the test toward humidity are illustrated in Figure 10 for OSR700 and SWP700 sensors. Both materials are sensitive toward humidity starting from low RH values. In fact, impedance begins to increase (i.e., SR% slope is negative) from 5 RH%, and an increment in impedance of

the film was noticed until about 40 RH% for OSR700 sensor and until 25 RH% for SWP700. In both cases, biochar materials are behaving as p-type semiconductors under low amounts of humidity. On the contrary, for higher RH values, the impedance decreased from about 40 RH% for OSR700 and from 25 RH% for SWP700. More in detail, for the more conductive OSR700 film, the initial impedance under dry air was equal to 163.9 k Ω , and the final impedance under 99 RH% dropped down to 9 k Ω , leading to a sensor response SR% equal to 94.5%, together with a maximum hysteresis of 52% under 59 RH% (Figure 10a). For SWP 700 film, due to the highest surface area and porosity compared to OSR700 biochar, a more consistent impedance variation was reported, and the impedance declined from 9.5 M Ω in dry condition until 222 k Ω in a humid environment, with a SR% of 97.7%, as reported in Table 4. This material exhibits a lower hysteresis, with a maximum value for this parameter of 23% under 55 RH% (Figure 10b).

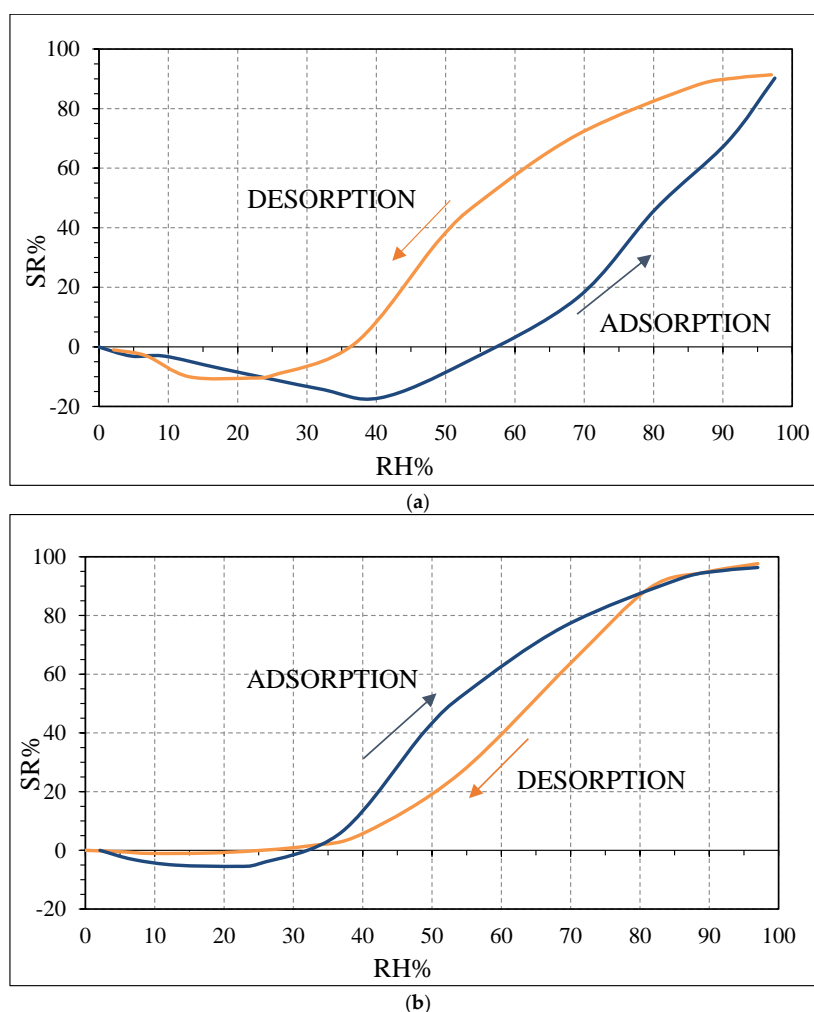


Figure 10. Sensor response towards relative humidity of (a) OSR700; (b) SWP700 biochars.

The response times, together with the recovery times were also determined. For OSR700 sensor they are 50 and 70 s respectively, while for SWP700 sensor both are in the order of 1 min.

Using SWP700, the biochar most sensitive towards humidity, sensors with different amounts of PVP were produced. PVP is a n-type hydrophilic conductive polymer [27,28] used here in 10 and 20 wt %. In addition, a pure PVP sensor was also realized. This reference sensor reported an impedance from 19 M Ω in dry condition to 420 k Ω under 98% of relative humidity with a response above 40 RH%. Results towards humidity of PVP sensor are shown in Figure 11 and confirm its n-type semiconducting behavior (the impedance decreases with the increasing RH values).

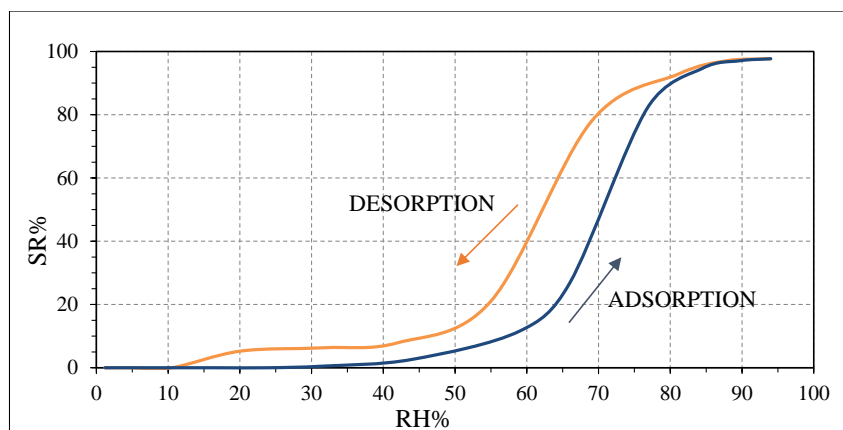


Figure 11. Polyvinylpyrrolidone (PVP) sensor response toward relative humidity.

From now on, the sample prepared with SWP700 biochar and 10 and 20 wt % PVP are labelled SWP700-10% PVP and SWP700-20% PVP, respectively. The sensors' response towards relative humidity are shown in Figure 12 and reported in Table 4. As expected, the sensitivity of the sensor decreased passing from the pure SWP700 biochar to the SWP700-10% PVP and SWP700-20% PVP sensors. Moreover, maximum hysteresis was equal to 42% under 60 RH% for SWP700-10% PVP composite, and 27% under 52 RH% for SWP700-20% PVP composite.

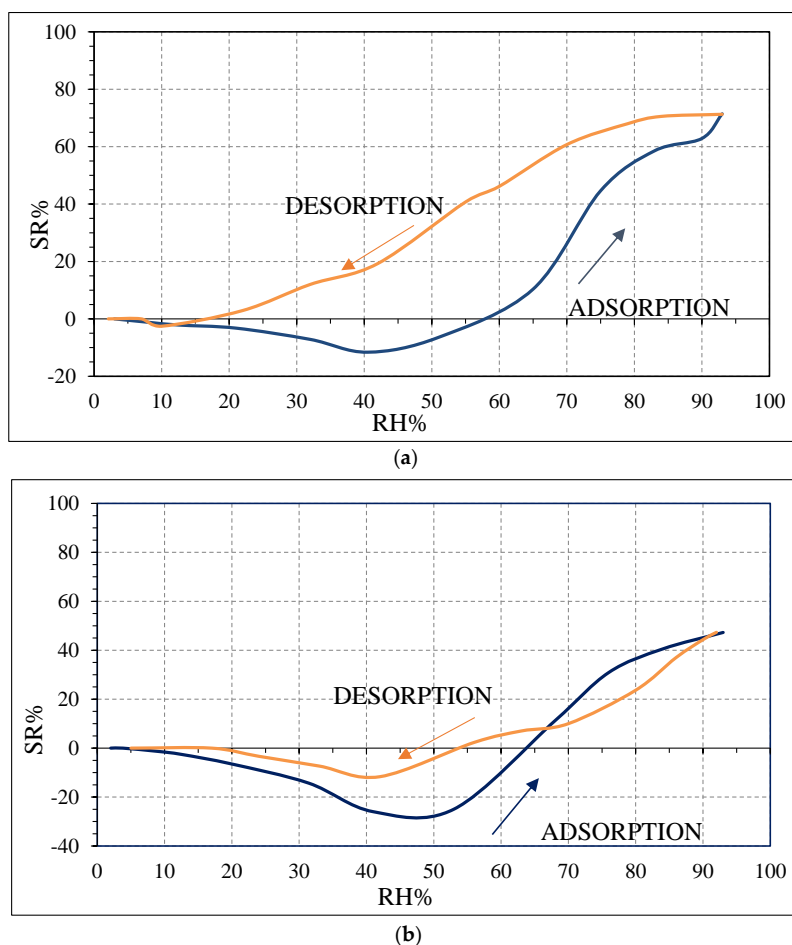


Figure 12. Sensor response toward relative humidity for SWP700 biochar with (a) 10%; (b) 20% of PVP acting as polymeric binder.

Table 4. Maximum SR% of PVP, OSR700, SWP700, SWP-10% PVP, and SWP-20% PVP sensors toward relative humidity at 26 °C.

Sensor	SR%
PVP	98
OSR700	94.5
SWP700	97.7
SWP700-10% PVP	65.2
SWP700-20% PVP	47.3

SWP700-10% PVP sensor seems to be a satisfying compromise between two fundamental parameters: sensitivity toward humidity and adhesion of the film onto the substrate. For this reason, evaluation of impedance's changes with the temperature together with cross sensitivity tests were performed for this sensor. The former results under a constant dry air flow of 1700 sccm are summarized in Table 5, where experiments were carried out in a thermostatic chamber between 25 °C and 60 °C.

Table 5. Impedance values of SWP700-10% PVP sensor at different temperature under dry air.

Temperature (°C)	Z ₀ (kΩ)
25	963
40	764
50	674
60	575

An increase in the operating temperature of 35 °C in dry conditions caused a drop in the film impedance of about 40%, as expected and reported in the literature [16,17,32].

Furthermore, cross sensitivity tests were performed under a constant flow of 1000 sccm on the same sensor, and the results are displayed in Table 6. No interferences were detected toward ozone 0.5 ppm, methane 100 ppm, and carbon dioxide 500 and 600 ppm, while only under 50 ppm of ammonia, an increase of 2.6% in impedance at room temperature was measured. Moreover, no synergic effect was observed for SWP700-10% PVP film under 600 ppm of carbon dioxide and air with 40 RH%.

Table 6. Impedance values of SWP700-10% PVP sensor toward ozone 0.5 ppm, ammonia 50 ppm, methane 100 ppm, and carbon dioxide 500 ppm.

Gas	Z ₀ (kΩ)	Z _g (kΩ)
O ₃ 0.5 ppm	954	954
NH ₃ 50 ppm	956	981
CH ₄ 100 ppm	958	959
CO ₂ 500 ppm	958	959
CO ₂ 600 ppm	958	959
RH40%	959	1130
CO ₂ 600 ppm + air RH40%	958	1120

4. Discussion

Water molecules can adsorb on the surface of carbon films due to a weak binding of water hydrogens to biochar surface carbon atoms [20]. Moreover, the sensing performance can be improved by the presence of defects in carbon grains, as evidenced by Raman spectroscopy results (Figure 5), which could create favorable adsorption sites to water molecules. These defects also favor water molecules adsorption by creating favorable adsorption sites [20]. Oxygen derivatives on carbon surface make the carbon surface hydrophilic and increase water vapor adsorption [20]. Thus, at low

RH values, a few water vapor molecules chemically adsorb on the biochar grain surfaces. Pati et al. [33] suggested a possible charge transfer between the adsorbate and the carbon film. This, probably leads to electrons exchange with a p-type semiconductor and their transfer to the valence band, leading to an increase of the impedance [20]. However, for higher RH values (above 40 RH% for OSR700 and above 25 RH% for SWP700), more water molecules are adsorbed on the surface, and operating at room temperature, water molecules will cluster to form a liquid-like multilayer of hydrogen-bonded water molecules [34]. Moreover, these physisorbed water molecules can condense into pores having a size in the range 1–250 nm. Since the formation of clusters of H₂O and the hydration of H⁺ into H₃O⁺ are energetically favored in liquid water, H⁺ are the dominant charge carriers in the water adsorbed in the mesopores. As the amount of H⁺ increases when increasing the moisture content, H⁺ can move freely in liquid water, leading to a decrease of grain surface impedance with increasing RH values [34]. This mechanism explains why Z_g , defined in Equation (1), tends to decrease with increasing RH values, leading to an enhancement of the sensor response, as observed in Figure 10, contrarily to what was expected from biochars as p-type semiconductors. Furthermore, the sp²/sp³ ratio determined by XPS measurements could also affect the sensor response at low relative humidity values (5–30%). In fact, the hydrophilic character is evidenced by a higher sp²/sp³ ratio, like in OSR700 film (sp²/sp³ = 0.45) compared to SWP700 one (sp²/sp³ = 0.06), in accordance with the literature [35]. As a consequence, the former biochar exhibits an enhanced affinity for the adsorption of water molecules at lower amounts of RH with respect to the latter one.

The difference in the hysteresis between the two biochars (Figure 10) may be probably explained by the different pore size distribution among the samples: in the case of OSR700, the hysteresis is higher because water molecules desorption is more difficult and takes longer time due to smaller pores or more tortuous ones. However, this feature was not investigated in this work and should be further studied. The effect of specific surface area on sensitivity of a quartz crystal microbalance-based polyacrylic acid ammonia sensor is reported in ref. [36]. The results of this study confirmed that the ammonia sensitivity was enhanced rapidly by the continuous increase of SSA and showed a logarithmic growth trend [36]. In our case, the higher SSA of SWP700 biochar probably favors water molecules adsorption too, leading to a faster decrease of the impedance value from 25 RH%, with respect to 40 RH% for OSR700 sensor.

Pristine PVP is a highly hydrophilic polymer due to the presence of a nitrogen atom in its molecule [37,38]. Thus, it can easily adsorb water molecules. As previously explained, at low relative humidity values, only a few water molecules can be adsorbed on the surface of PVP to generate a few H⁺ and H₃O⁺ ions. Then, the intrinsic electrons of PVP are the main responsible for the conduction of PVP which behaves as a n-type semiconductor as confirmed in Figure 11 [37,38]. When increasing RH values, the water molecules on the surface of PVP provide more carriers (H⁺ and H₃O⁺) which mainly contribute to the conduction of PVP [39]. Above 94 RH%, the pristine PVP can be swelled by water, generating many conductive ions [40].

The decrease in impedance for the SWP700-10% PVP sensor compared to the SWP700 is probably due to the enhanced connection between biochar grains in the polymeric medium compared to biochar as it is. Moreover, when n-p heterojunctions are formed at nanoscale level between two semiconductors, i.e., PVP and biochar, electrons are transferred from n-type PVP to p-type biochar, while holes are moved in the opposite direction, to equate the Fermi levels of both materials [41]. Then, a potential barrier with the bending of energy levels is formed at the heterojunctions. Thus, the impedance of the n-p composite increases, compared to the pure biochar [41]. With increasing initial impedance, the variation of impedance due to the presence of few water molecules becomes more evident, leading to a higher sensitivity, as experimentally observed for the sensors SWP700-10% PVP and SWP700-20% PVP (Figure 12a,b), compared to the sensor SWP700 (Figure 10b). This larger initial impedance leads to a stronger initial response [41]. Table 7 compares the results presented in this paper and literature data when possible because the initial resistance value under dry air is not always mentioned or determinable from experimental curves. The obtained results are rather promising.

Table 7. Response toward humidity of carbon-based and of composite polymer-based resistive sensors at room temperature.

Sensing Material and Preparation Technique	Sensor Response	Reference
Carbon thin-film made by spray pyrolysis technique	$R_g/R_0 \approx 0.95$ under 2 RH%	[16]
Pyrolyzed medium density fiberboard impregnated with phenolic resin	Resistance linearly changed in the 10–70 RH% range	[17]
High frequency PVD carbon thin-films	$R_g/R_0 \approx 1.08$ from 11 RH%	[20]
Pyrolyzed bamboo screen-printed thick-films	$R_g/R_0 = 0.98$ under 10 RH%	[26]
Spin-coated thin-films of MWCNTs/PMMA doped with KOH	The log-impedance curve linearly changed in the 30–90 RH% range from 10^4 to $10^7 \Omega$	[42]
Spin-coated PEDOT: PSS/iron oxide nanoparticle composite thin-films	Resistance linearly changed in the 30–70 RH% range	[43]
Drop casting of MWCNTS/PAA composite films	Almost linear resistance change in the range 30–90 RH%	[44]
Drop casting of N-doped carbon sphere-poly (vinyl alcohol) composites	Conductance (G) determined from 9 RH%. $S = \frac{\Delta \log G}{\Delta(RH\%)} = 17$	[45]
Pellet of tin oxide nanoparticles encapsulated in polyaniline matrix	$R_g/R_0 = 0.98$ under 10 RH%	[46]
Drop-coating biochar/PVP	R_g/R_0 1.02 = under 10 RH%	This work

(NB: PVD = physical vapor deposition; MWCNTs = multi-walled carbon nano-tubes; PMMA = poly(methylmetacrylate); PEDOT = poly(3,4-ethylenedioxythiophene); PSS = polystyrene sulfonate; PAA = poly (acrylic acid); RGO = reduced graphene oxide).

Future prospective involves monitoring sensors' response in function of the temperature, since a preliminary evaluation of carbonaceous materials change resistance values with temperature was performed only under dry air. Moreover, ageing phenomena will be investigated too, and verifying how their response is eventually affected by storage must be investigated.

5. Conclusions

In this work, two commercial biochars were evaluated as new humidity sensing materials.

The sensors were prepared by a simple and low-cost process, and their humidity sensing properties were investigated at room temperature in the relative humidity range from 0.0 to almost 100%. Both pristine materials are sensitive toward humidity starting from low RH values. In fact, impedance begins to increase from 5 RH%, and an increment in impedance was noticed until about 40 RH% for the OSR700 sensor, and until 25 RH% for the SWP700 sensor. In both cases, biochar materials behave as p-type semiconductors under low amounts of humidity. On the contrary, for higher RH values, the impedance decreased due to a higher amount of water molecules adsorption.

When PVP is added as a polymeric binder to SWP700 biochar, n-p heterojunctions are formed between the two semiconductors, leading to a higher sensitivity at low RH values for the sensors SWP700-10% PVP and SWP700-20% PVP with respect to the pure SWP700 sensor. This behavior makes these sensors potentially interesting for industrial applications where low RH values are required (drying of clothes (0–40 RH%), pharmaceuticals (20–40 RH%), cereal stocking (0–45 RH%), electronic parts, ceramic powders, humidity control in factories, dryers, dried foodstuffs (0–50 RH%), and film desiccation (0–30 RH%) . . .) [34]. Thus, they are rather devoted to niche markets rather than the mass market.

Finally, the response and recovery times were reasonably fast (in the order of 1 min), and the cross-sensitivity to methane, carbon dioxide, ammonia, and ozone was negligible.

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